Claims

We claim:

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1. A compound of the structure

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$$R^{1}$$
 G Q R^{2} R^{3} R^{4} R^{4}

wherein A is selected from the group consisting of O, S, and NR⁵;

E is selected from the group consisting of CH₂, O, S, and NR6;

Q is selected from the group consisting of C(O) and (CH₂)_k wherein k is an integer of 0 or 1;

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J is selected from the group consisting of O, S and NR⁸;

G is selected from the group consisting of O, NH, S, and (CH₂)_p wherein p is an integer of 0 or 1;

T is selected from the group consisting of C(O) and (CH₂)_b wherein b is an integer of from 0 to 3;

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L is selected from the group consisting of O, NR⁷, S, and

 $(CH_2)_n$ wherein n is an integer of 0 or 1;

M is selected from the group consisting of $C(R^9)(R^{10})$ and (CH₂), wherein u is an integer of from 0 to 3;

X is selected from the group consisting of CO₂B, PO₃H₂, SO₃H, OPO₃H₂, C(O)NHC(O)R¹¹, C(O)NHSO₂R¹², tetrazolyl and hydrogen;

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B, R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹ and R¹² are independently selected from the group consisting of hydrogen, alkyl, cycloalkyl,

aryl, hydroxyalkyl, alkoxy, alkoxyalkoxy, cycloalkylalkyl, alkylamino, haloalkyl, alkylaryl, arylalkyl, heterocyclyl, alkylheterocyclyl and heterocyclylalkyl groups;

wherein R^2 and R^3 taken together may form a ring; R^4 and R^7 taken together may form a ring; R^9 and R^{10} taken together may form a ring; and salts thereof.

2. A compound of claim 1 wherein

R¹, R² and R³ are independently selected from the group consisting of hydrogen, alkoxy, alkoxyalkoxy, aryl, alkylaryl, arylalkyl, heterocyclyl and alkyl;

R⁴ is selected from the group consisting of aryl, alkylaryl, arylalkyl, heterocyclyl, alkylheterocyclyl and heterocyclylalkyl;

X is CO₂B; and

M is C(R⁹)(R¹⁰) wherein R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen and lower alkyl.

- 3. A compound of claim 1 further comprising derivatives of said compound selected from the group consisting of esters, carbamates, aminals, amides, and pro-drugs thereof.
- 4. A compound of claim 1 of the structure

wherein A is selected from the group consisting of O, S, and NR⁵;
E is selected from the group consisting of CH₂, O, S, and
NR⁶;

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Q is selected from the group consisting of $C(O)$ and $(CH_2)_k$ wherein k
is an integer of 0 or 1;
G is selected from the group consisting of O, NH, S, and (CH ₂) _p
wherein p is an integer of 0 or 1;
T is selected from the group consisting of C(O) and (CH ₂) _b wherein b is
an integer of 0 to 3;
L is selected from the group consisting of O, NR ⁷ , S, and
(CH ₂) _n wherein n is an integer of 0 or 1;
B, R ¹ , R ² , R ³ , R ⁴ , R ⁵ , R ⁶ , R ⁷ , R ⁹ and R ¹⁰ are independently selected from
the group consisting of hydrogen, alkyl, cycloalkyl, aryl,
hydroxyalkyl, alkoxy, alkoxyalkoxy, cycloalkylalkyl,
alkylamino, haloalkyl, alkylaryl, arylalkyl, heterocyclyl,
alkylheterocyclyl and heterocyclylalkyl groups;
wherein R ² and R ³ taken together may form a ring;
R ⁴ and R ⁷ taken together may form a ring;
R ⁹ and R ¹⁰ taken together may form a ring;
and salts thereof.
5. A compound of claim 4 wherein R ¹ , R ² and R ³ are independently
selected from the group consisting of hydrogen, alkoxy,
alkoxyalkoxy, aryl, alkylaryl, arylalkyl, heterocyclyl and alkyl;
R ⁴ is selected from the group consisting of aryl, alkylaryl, arylalkyl,
heterocyclyl, heterocyclylalkyl and alkyheterocyclyl;
R ⁵ and R ⁶ are hydrogen; and
R ⁹ and R ¹⁰ are independently selected from the group consisting of
hydrogen and lower alkyl.

A compound of claim 4 further comprising derivatives of said

compound selected from the group consisting of esters, carbamates, aminals, amides, and pro-drugs thereof.

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7. A compound of claim 1 selected from the group consisting of: (3S)-3-(1,3-benzodioxol-5-yl)-3-((((1S)-3-(methylsulfanyl)-1-((phenylsulfanyl)methyl)propyl)amino) carbonyl)amino)propanoic acid, (3S)-3-(1,3-benzodioxol-5-yl)-3-(((((1S)-2-((cyclopropylmethyl)thio)-1-5 ((phenylthio)methyl)ethyl)amino)carbonyl) amino)propanoic acid, (9S,13S)-13-(1,3-benzodioxol-5-yl)-3,11-dioxo-1-phenyl-9-{[(2thienylmethyl)amino|carbonyl}-2-oxa-4,10,12-triazapentadecan-15-oic acid, (9S,13S)-13-(1,3-benzodioxol-5-yl)-9-{[(3-hydroxy-4methoxybenzyl)amino]carbonyl}-3,11-dioxo-2-oxa-4,10,12-10 triazapentadecan-15-oic acid, $(3S)-3-(1,3-benzodioxol-5-yl)-3-\{[({(1S)-2-(benzylsulfanyl)-1-}$ [(phenylsulfanyl)methyl]ethyl}amino)carbonyl] amino) propanoic acid, (3S)-3-(1,3-benzodioxol-5-yl)-3- $\{[(\{(1S)$ -3-(methylsulfanyl)-1-[({4-[(2-toluidinocarbonyl)amino]phenyl}sulfanyl) methyl|propyl}amino)carbonyl|amino}propanoic acid, (3S)-3-(1,3-15 benzodioxol-5-yl)-3- $\{[(\{(1S)-2-(ethylsulfanyl)-1-(ethylsulfanyl)$ [(phenylsulfanyl)methyl]ethyl}amino) carbonyl]amino}propanoic acid, (9S,13S)-13-(1,3-benzodioxol-5-yl)-9-[({4-[(2methylbenzyl)amino|benzyl} amino)carbonyl]-3,11-dioxo-1-phenyl-2-20 oxa-4,10,12-triazapentadecan-15-oic acid, (3S)-3-(1,3-benzodioxol-5yl)-3-{[({(1S)-3-(methylsulfanyl)-1-[({3-[(2toluidinocarbonyl)amino[phenyl] sulfanyl)methyl] propyl\amino\carbonyl\amino\propanoic acid, (3S)-3-(1,3benzodioxol-5-yl)-3-{[({(1S)-2-(ethylthio)-1-25 [(phenylthio)methyl]ethyl}oxy)carbonyl]amino} propanoic acid, (9S, 13S)-13-(1,3-benxodioxol-5-yl)-3,11-dioxo-1-phenyl-9-(((4-((2toluidinocarbonyl)amino)benzyl)amino)carbonyl)-2-oxa-4, 10,12triazapentadecan-15-oic acid, and pharmaceutically acceptable salts thereof.

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- 8. A compound of claim 7 further comprising derivatives of said compound selected from the group consisting of esters, carbamates, aminals, and amides, optical isomers and pro-drugs thereof.
- 9. A pharmaceutical composition comprising:
 a compound of claim 1
 and pharmaceutically acceptable salts thereof,
 in a pharmaceutically acceptable carrier.
- 10. A method for selectively inhibiting $\alpha_4\beta_1$ integrin binding in a mammal comprising administering to said mammal a therapeutic amount of a compound of claim 1.

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